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NEWS 1
                Web Page URLs for STN Seminar Schedule - N. America
NEWS 2
                "Ask CAS" for self-help around the clock
NEWS 3 JAN 17 Pre-1988 INPI data added to MARPAT
NEWS 4 FEB 21 STN AnaVist, Version 1.1, lets you share your STN AnaVist
                visualization results
NEWS 5 FEB 22 The IPC thesaurus added to additional patent databases on STN
NEWS 6 FEB 22 Updates in EPFULL; IPC 8 enhancements added
NEWS 7 FEB 27 New STN AnaVist pricing effective March 1, 2006
NEWS 8 MAR 03 Updates in PATDPA; addition of IPC 8 data without attributes
NEWS 9 MAR 22 EMBASE is now updated on a daily basis
NEWS 10 APR 03 New IPC 8 fields and IPC thesaurus added to PATDPAFULL
NEWS 11 APR 03
                Bibliographic data updates resume; new IPC 8 fields and IPC
                thesaurus added in PCTFULL
NEWS 12 APR 04
                STN AnaVist $500 visualization usage credit offered
NEWS 13 APR 12 LINSPEC, learning database for INSPEC, reloaded and enhanced
NEWS 14 APR 12 Improved structure highlighting in FQHIT and QHIT display
                in MARPAT
NEWS 15 APR 12 Derwent World Patents Index to be reloaded and enhanced during
                second quarter; strategies may be affected
NEWS 16 MAY 10 CA/CAplus enhanced with 1900-1906 U.S. patent records
NEWS 17 MAY 11 KOREAPAT updates resume
NEWS 18 MAY 19 Derwent World Patents Index to be reloaded and enhanced
NEWS EXPRESS
            FEBRUARY 15 CURRENT VERSION FOR WINDOWS IS V8.01a,
             CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
```

CURRENT MACINTOSH VERSION FOR WINDOWS IS V8.01a,
CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005.
V8.0 AND V8.01 USERS CAN OBTAIN THE UPGRADE TO V8.01a AT
http://download.cas.org/express/v8.0-Discover/

NEWS HOURS STN Operating Hours Plus Help Desk Availability
NEWS LOGIN Welcome Banner and News Items
NEWS IPC8 For general information regarding STN implementation of IPC 8
NEWS X25 X.25 communication option no longer available after June 2006

Enter NEWS followed by the item number or name to see news on that specific topic.

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If you provide us with your name, login ID, and e-mail address, you will be entered in a drawing to win a free iPod(R). Your responses will be kept confidential and will help us make future improvements to STN.

Take survey: http://www.zoomerang.com/survey.zgi?p=WEB2259HNKWTUW

Thank you in advance for your participation.

FILE 'HOME' ENTERED AT 14:55:08 ON 24 MAY 2006

=>

Uploading

THIS COMMAND NOT AVAILABLE IN THE CURRENT FILE Do you want to switch to the Registry File? Choice (Y/n):

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Some commands only work in certain files. For example, the EXPAND command can only be used to look at the index in a file which has an index. Enter "HELP COMMANDS" at an arrow prompt (=>) for a list of commands which can be used in this file.

=> FILE REGISTRY

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

FILE 'REGISTRY' ENTERED AT 14:55:20 ON 24 MAY 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 American Chemical Society (ACS)

Property values tagged with IC are from the $\dot{\text{ZIC/VINITI}}$ data file provided by InfoChem.

STRUCTURE FILE UPDATES: 23 MAY 2006 HIGHEST RN 885357-09-5 DICTIONARY FILE UPDATES: 23 MAY 2006 HIGHEST RN 885357-09-5

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH January 6, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

* The CA roles and document type information have been removed from * the IDE default display format and the ED field has been added, * effective March 20, 2005. A new display format, IDERL, is now * available and contains the CA role and document type information. *

Structure search iteration limits have been increased. See HELP SLIMITS for details.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/ONLINE/UG/regprops.html

=>

Uploading C:\Program Files\Stnexp\Queries\10635294a.str

chain nodes : 12 13 14 15 16 17 ring nodes : 5 6 7 8 10 11 chain bonds : 1-12 2-13 8-12 12-17 13-14 13-15 14-16 ring bonds : 1-2 1-5 2-3 3-4 4-5 6-7 6-11 7-8 8-9 9-10 10-11 exact/norm bonds : 1-2 1-5 1-12 3-4 4-5 12-17 13-14 13-15 14-16 exact bonds : 2-3 2-13 8-12 normalized bonds : 6-7 6-11 7-8 8-9 9-10 10-11 isolated ring systems : containing 1 : 6 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS 19:CLASS

10635294c.trn

Stereo Bonds:

17-12 (Single Wedge).

Stereo Chiral Centers:

12 (Parity=Don't Care)

Stereo RSS Sets:

Type=Relative (Default). 1 Nodes= 12

L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR

Structure attributes must be viewed using STN Express query preparation.

=> s 11

SAMPLE SEARCH INITIATED 14:55:34 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 468 TO ITERATE

100.0% PROCESSED 468 ITERATIONS

1 ANSWERS

26 MSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

PROJECTED ITERATIONS: 8063 TO 106.

PROJECTED ITERATIONS: 8063 TO 10657 PROJECTED ANSWERS: 1 TO 80

L2 1 SEA SSS SAM L1

=> s l1 sss full

FULL SEARCH INITIATED 14:55:40 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 9425 TO ITERATE

100.0% PROCESSED 9425 ITERATIONS

10635294c.trn Page 4 14:59

SEARCH TIME: 00.00.01

L3 26 SEA SSS FUL L1

=> FIL HCAPLUS

COST IN U.S. DOLLARS SINCE FILE TOTAL

FULL ESTIMATED COST ENTRY SESSION 166.94 167.15

FILE 'HCAPLUS' ENTERED AT 14:55:46 ON 24 MAY 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

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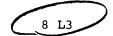
FILE COVERS 1907 - 24 May 2006 VOL 144 ISS 22 FILE LAST UPDATED: 23 May 2006 (20060523/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 13 L4

L5



=> s 14 and radioactive halogen

173123 RADIOACTIVE

28 RADIOACTIVES

173132 RADIOACTIVE

(RADIOACTIVE OR RADIOACTIVES)

106628 HALOGEN

21560 HALOGENS

117523 HALOGEN

(HALOGEN OR HALOGENS)

78 RADIOACTIVE HALOGEN

(RADIOACTIVE (W) HALOGEN)

1 L4 AND RADIOACTIVE HALOGEN

=> s l4 and radioactive

173123 RADIOACTIVE

28 RADIOACTIVES

173132 RADIOACTIVE

(RADIOACTIVE OR RADIOACTIVES)

L6 2 L4 AND RADIOACTIVE

=> d 15 ibib abs hitstr tot

L5 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2006 ACS on STN

10635294c.trn

Page 5

14:59

ACCESSION NUMBER: 2005:123220 HCAPEÚS

DOCUMENT NUMBER: 142:198079

Preparation of radiolabeled 1-(phenylethyl)imidazole-5-carboxylc acid ester derivatives TITLE:

Zolle, Mse; Hammerschmidt, Friedrich

mile

PATENT ASSIGNEE(S): Austria

SOURCE: U.S. Pat. Appl. Publ., 15 pp.

CODEN: USXXCO

DOCUMENT TYPE:

Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

INVENTOR (S):

PATENT NO. KIND DATE APPLICATION NO. DATE US 2005033060 A1 20050210 US 2003-635294 20030806 US 2003-635294 PRIORITY APPLN. INFO.: 20030806

OTHER SOURCE(S): CASREACT 142:198079; MARPAT 142:198079

GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Halogenated carboxylic ester derivs. of phenylethylimidazole (I) [R1 = linear or branched C1-4 alkyl which is optionally substituted with a halogen selected from the groups consisting of F, Cl, I or Br; R2 = C1-2 alkyl; X = a nonradioactive or a radioactive halogen] or (II) [X = a nonradioactive or radioactive halogen selected from the group consisting of I, Br, and F; X = aradioactive halogen selected from the group consisting of 123I, 124I, 131I, 76Br, 82Br or 18F] are prepared via coupling of (S)-secondary alc. (III) (R2, X = same as above) with imidazolecarboxylate ester (IV) (R1 = same as above). Radio-halogenated forms of these compds. are ideally suited for positron-imaging of the adrenal glands, as it is known that these compds. demonstrate a selective and high rate of accumulation in the adrenals. The method of preparing these derivs. proceeds by the conversion of a stable, non-radioactive intermediate having trialkylstannyl leaving groups (V) [R1, R2 = same as above; L = an alkylstannyl group selected from the group consisting of trimethylstannyl, triethylstannyl, tri-n-propylstannyl and tri-n-butylstannyl] and (VI) (R1, R2 = same as above). These intermediates are efficiently converted to the corresponding halogenated forms by substitution of the trialkylstannyl group with the halogen or radiohalogen. Thus, 4-iodoacetophenone was reduced by DIBAH in toluene/Et2O at -78° to give 86% 1-(4-iodophenyl)ethanol which was esterified by chloroacetic anhydride in the presence of pyridine in CH2Cl2 at 0° for 2 h to give 91% 1-(4-iodophenyl)ethyl chloroacetate (VII). VII underwent enzymic hydrolysis in the presence of lipase SAM II in a mixture of tert-Bu Me ether and phosphate buffer at 0° for 2 h while keeping pH at 7.0 by adding 0.5 N aqueous NaOH solution to give 43% (R)-1-(4-iodophenyl)ethanol (98% ee) and 44% (S)-1-(4-iodophenyl)ethyl chloroacetate (>98% ee) (VIII). VIII was coupled with Me 3H-imidazole-4-carboxylate using triphenylphosphine and di(tert-butyl) azocarboxylate in THF at -30° to 0° over 2 .5 h to give 67% (R)-(+)-Me 3-[1-(4-iodophenyl)ethyl]-3H-imidazole-4-carboxylate (99% ee) which was refluxed with hexamethyltin in toluene at 135° for 17 h to give 96% (R)-(+)-Me 3-[1-[4-(trimethylstannyl)phenyl]ethyl]-3H-imidazole-4-carboxylate (IX).

IX (30 μg) was reacted with [131I]iodide in 10-20 μL 0.05 N aqueous NaOH solution, 15 μL aqueous chloramine-T solution (1 mg/mL), and 6 μL 1 N aqueous HCl

solution at room temperature for 1 min to give (R) - (+) - Me 3 - [1 - (4 -[131I]iodophenyl)ethyl]-3H-imidazole-4-carboxylate (131I-MTO), i.e. II (R1 = R2 = Me, X = 131I).

IT 813466-09-0P

> RL: BSU (Biological study, unclassified); DGN (Diagnostic use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES

(preparation of radiolabeled (phenylethyl)imidazolecaboxylic acid ester derivs. as positron-emission imaging agents for adrenal glands)

RN 813466-09-0 HCAPLUS

CN 1H-Imidazole-5-carboxylic acid, 1-[(1R)-1-[4-(iodo-131I)phenyl]ethyl]-, methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).

IT 813466-05-6P, (R)-(+)-Methyl 3-[1-(4-Iodophenyl)ethyl]-3Himidazole-4-carboxylate

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of radiolabeled (phenylethyl)imidazolecaboxylic acid ester derivs. as positron-emission imaging agents for adrenal glands)

RN 813466-05-6 HCAPLUS

1H-Imidazole-5-carboxylic acid, 1-[(1R)-1-(4-iodophenyl)ethyl]-, methyl CN ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).

=> d l6 ibib abs hitstr tot

ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:123220 HCAPLUS

DOCUMENT NUMBER: 142:198079

TITLE: Preparation of radiolabeled 1-(phenylethyl)imidazole-5-

carbox lic acid ester derivatives (Zolle, Ilse; Hammerschmidt, Friedrich

INVENTOR (S): PATENT ASSIGNEE(S): Austri

10635294c.trn Page 7

14:59

SOURCE: U.S. Pat. Appl. Publ., 15 pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

US 2005033060 A1 20050210 US 2003-635294 20030806

PRIORITY APPLN. INFO: US 2003-635294 20030806

OTHER SOURCE(S): CASREACT 142:198079; MARPAT 142:198079

GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Halogenated carboxylic ester derivs. of phenylethylimidazole (I) [R1 = linear or branched C1-4 alkyl which is optionally substituted with a halogen selected from the groups consisting of F, Cl, I or Br; R2 = C1-2alkyl; X = a nonradioactive or a radioactive halogen] or (II) [X = a nonradioactive or radioactive halogen selected from the group consisting of I, Br, and F; X = a radioactive halogen selected from the group consisting of 123I, 124I, 131I, 76Br, 82Br or 18F] are prepared via coupling of (S)-secondary alc. (III) $(R2, X = same \ as$ above) with imidazolecarboxylate ester (IV) (R1 = same as above). Radio-halogenated forms of these compds. are ideally suited for positron-imaging of the adrenal glands, as it is known that these compds. demonstrate a selective and high rate of accumulation in the adrenals. The method of preparing these derivs. proceeds by the conversion of a stable, non-radioactive intermediate having trialkylstannyl leaving groups (V) [R1, R2 = same as above; L = an alkylstannyl group selected from the group consisting of trimethylstannyl, triethylstannyl, tri-n-propylstannyl and tri-n-butylstannyl] and (VI) (R1, R2 = same as above). These intermediates are efficiently converted to the corresponding halogenated forms by substitution of the trialkylstannyl group with the halogen or radiohalogen. Thus, 4-iodoacetophenone was reduced by DIBAH in toluene/Et2O at -78° to give 86% 1-(4-iodophenyl)ethanol which was esterified by chloroacetic anhydride in the presence of pyridine in CH2Cl2 at 0° for 2 h to give 91% 1-(4-iodophenyl)ethyl chloroacetate (VII). VII underwent enzymic hydrolysis in the presence of lipase SAM II in a mixture of tert-Bu Me ether and phosphate buffer at 0° for 2 h while keeping pH at 7.0 by adding 0.5 N aqueous NaOH solution to give 43% (R)-1-(4-iodophenyl)ethanol (98% ee) and 44% (S)-1-(4-iodophenyl)ethyl chloroacetate (>98% ee) (VIII). VIII was coupled with Me 3H-imidazole-4-carboxylate using triphenylphosphine and di(tert-butyl) azocarboxylate in THF at -30° to 0° over 2 .5 h to give 67% (R)-(+)-Me 3-[1-(4-iodophenyl)]-3H-imidazole-4-carboxylate (99% ee) which was refluxed with hexamethyltin in toluene at 135° for 17 h to give 96% (R)-(+)-Me 3-[1-[4-(trimethylstannyl)phenyl]ethyl]-3H-imidazole-4-carboxylate (IX). IX (30 μg) was reacted with [131I]iodide in 10-20 μL 0.05 N aqueous NaOH solution, 15 μL aqueous chloramine-T solution (1 mg/mL), and 6 μL 1 N aqueous HCl solution at room temperature for 1 min to give $(R) - (+) - Me \ 3 - [1 - (4 -$ [1311]iodophenyl)ethyl]-3H-imidazole-4-carboxylate (1311-MTO), i.e. II (R1 $= R2 = Me, \dot{X} = 131I).$

IT 813466-09-0P

RL: BSU (Biological study, unclassified); DGN (Diagnostic use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of radiolabeled (phenylethyl)imidazolecaboxylic acid ester derivs. as positron-emission imaging agents for adrenal glands)

RN 813466-09-0 HCAPLUS

CN 1H-Imidazole-5-carboxylic acid, 1-[(1R)-1-[4-(iodo-131I)phenyl]ethyl]-, methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).

IT 813466-05-6P, (R)-(+)-Methyl 3-[1-(4-Iodophenyl)ethyl]-3H-

imidazole-4-carboxylate

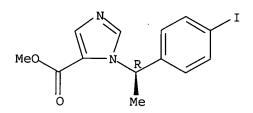
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of radiolabeled (phenylethyl)imidazolecaboxylic acid ester derivs. as positron-emission imaging agents for adrenal glands)

RN 813466-05-6 HCAPLUS

CN 1H-Imidazole-5-carboxylic acid, 1-[(1R)-1-(4-iodophenyl)ethyl]-, methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).



L6 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1988:167377 HCAPLUS

DOCUMENT NUMBER: 108:167377

TITLE: Synthesis of (R)-(+)-3H-etomidate

AUTHOR(S): Janssen, Cor G. M.; Thijssen, Jos B. A.; Verluyten,

Willy L. M.; Heykants, Jozef J. P.

CORPORATE SOURCE: Dep. Drug Metab. Pharmacokinet., Janssen Pharm.,

Beerse, B-2340, Belg.

SOURCE: Journal of Labelled Compounds and Radiopharmaceuticals

(1987), 24(8), 909-18

CODEN: JLCRD4; ISSN: 0362-4803

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 108:167377

GI

AB Etomidate, (R)-(+)-ethyl-1-(1-phenylethyl)-1H-imidazole-5-carboxylate (I, R = H) is a short-acting hypnotic. A new synthesis, featuring optical resolution on a non-radioactive precursor and introduction of the tritium label by reductive dehalogenation of I (R = Br) is described. I (R = T) was obtained at a specific activity of 3.77 Ci/mmol and a 99.9% purity.

IT 112366-36-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and desulfurization of, with sodium nitrite)

RN 112366-36-6 HCAPLUS

CN lH-Imidazole-4-carboxylic acid, 3-[1-(2-bromophenyl)ethyl]-2,3-dihydro-2-thioxo-, ethyl ester, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 112366-50-4P

RN 112366-50-4 HCAPLUS

CN 1H-Imidazole-5-carboxylic acid, 1-[1-(2-bromophenyl)ethyl]-, ethyl ester, (R)-, sulfate (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 112366-49-1

CMF C14 H15 Br N2 O2

Absolute stereochemistry.

CM 2

CRN 7664-93-9 CMF H2 O4 S

IT 112366-49-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation, reductive debromination, and tritiation of)

RN 112366-49-1 HCAPLUS

CN 1H-Imidazole-5-carboxylic acid, 1-[1-(2-bromophenyl)ethyl]-, ethyl ester, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

=> log y COST IN U.S. DOLLARS

COST IN U.S. DOLLARS
SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)
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CA SUBSCRIBER PRICE ENTRY SESSION -2.25 -2.25

STN INTERNATIONAL LOGOFF AT 14:57:33 ON 24 MAY 2006